FDA 510(k) Premarket Notification

CLEARED

K831857 SURGITEK PERCUTANEOUS ARNEGRADE SILITEK

Aug 11, 1983
63 days to decision

K831857 · Product code: **FAD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k831857/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Jun 9, 1983
Decision date	Aug 11, 1983
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Medical Engineering Corp.**
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-1993